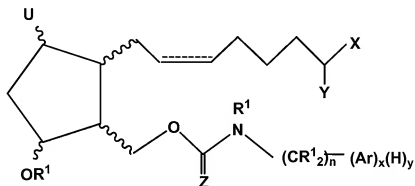


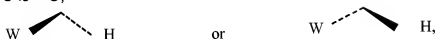
LISTING OF THE CLAIMS

1. (Currently Amended) A method of treating ocular hypertension which comprises administering to a mammal having ocular hypertension a therapeutically effective amount of a compound represented by formula I:



wherein a wavy segments indicate either the \square or \square configuration; the dashed bond represents a double bond or a single bond;

U is $=O$;

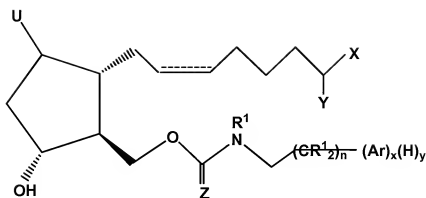


wherein W is halogen;

Z is O or S;

Ar is selected from the group consisting of aryl or heteroaryl radicals having from 4 to 10 carbon atoms and substituted derivatives of said aryl and heteroaryl radicals; n is 0 or an integer of from 1 to 4; x and y are 1 or 0, provided that when x is 1, y is 0 and when x is 0, y is 1; R¹ is hydrogen or a lower alkyl radical or a substituted lower alkyl radical having up to six carbon atoms; X is selected from the group consisting of -OR¹ and -N(R¹)₂; Y is =O or represents 2 hydrogen radicals, Z is S or O; wherein the substituent on the lower alkyl, aryl or heteroaryl radical is selected from the group consisting of lower alkyl, hydroxy, lower alkyloxy, halogen, trifluoromethyl (CF₃), COR₁, COCF₃, SO₂NR₁, SO₂NH₂, NO₂ and CN and/or the pharmaceutically acceptable salts of said compounds and/or esters.

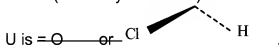
2. (Original) The method of claim 1 wherein said compound is represented by formula II:



wherein n is 0 or 1, 2, 3 or 4; hatched lines at position C-8 and C-11 indicate the α orientation; and the triangle at position C-12 represents the β orientation.

3. (Original) The method of claim 2 wherein Y is = O and X is $-OR^1$.

4. (Currently Amended) The method of claim 3 wherein



5. (Original) The method of claim 4 wherein Z is O.

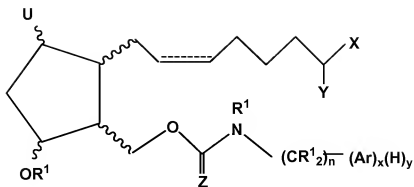
6. (Original) The method of claim 4 wherein R^1 is H or methyl.

7. (Original) The method of claim 4 wherein Ar is phenyl.

8. (Original) The method of claim 4 wherein x is 0.

9. (Original) An ophthalmic solution comprising a therapeutically effective amount of a compound of formula I, as defined in Claim 1, or a pharmaceutically acceptable salt thereof, in admixture with a non-toxic, ophthalmically acceptable liquid vehicle, packaged in a container suitable for metered application.

10. (Currently Amended) The ophthalmic solution of Claim 9 wherein said compound is a compound of Formula III:



wherein a wavy segments indicate either the \square or \square configuration; the dashed bond represents a double bond or a single bond;

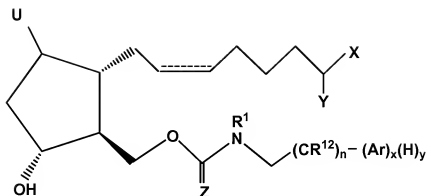


wherein W is halogen;

Z is O or S;

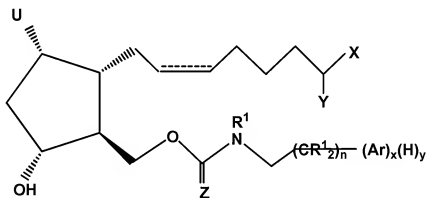
Ar is selected from the group consisting of aryl or heteroaryl radicals having from 4 to 10 carbon atoms and substituted derivatives of said aryl and heteroaryl radicals; n is 0 or an integer of from 1 to 4; x and y are 1 or 0, provided that when x is 1, y is 0 and when x is 0, y is 1; R¹ is hydrogen or a lower alkyl radical or a substituted lower alkyl radical having up to six carbon atoms; X is selected from the group consisting of -OR¹ and -N(R¹)₂; Y is =O or represents 2 hydrogen radicals; wherein the substituent Z is S or O; wherein the substituent on the lower alkyl, aryl or heteroaryl radical is selected from the group consisting of lower alkyl, hydroxy, lower alkyloxy, halogen, trifluoromethyl (CF₃), COR₁, COCF₃, SO₂NR₁, SO₂NH₂, NO₂ and CN and/or the pharmaceutically acceptable salts of said compounds and/or esters.

14. (Original) The compound of claim 13 wherein said compound is formula II:



wherein n is 0 or 1, 2, 3 or 4; hatched lines at position C-8 and C-11 indicate the α orientation; and the triangle at position C-12 represents the β orientation.

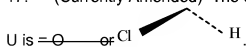
15. (Original) The compound of claim 14 wherein said compound is represented by formula II:



wherein n is 0 or 1, 2 or 4; hatched lines at position C-8 and C-11 indicate the α orientation; and the triangle at position C-12 represents the β orientation.

16. (Original) The compound of claim 15 wherein Y is $=O$ and X is $-OR^1$.

17. (Currently Amended) The compound of claim 16 wherein



18. (Original) The compound of claim 17 wherein Z is O .

19. (Original) The compound of claim 18 wherein R¹ is H or methyl.
20. (Original) The compound of claim 19 wherein Ar is phenyl.
21. (Original) The method of claim 1 wherein said compound is selected from the group consisting of

(Z)-7-((1R,2S,3R)-2-Butylcarbamoyloxymethyl-3-hydroxy-5-oxo-cyclopentyl)-hept-5-enoic acid methyl ester

(Z)-7-((1R,2S,3R)-2-Butylcarbamoyloxymethyl-3-hydroxy-5-oxo-cyclopentyl)-hept-5-enoic acid

(Z)-7-((1R,2S,3R,5R)-2-Butylcarbamoyloxymethyl-5-chloro-3-hydroxy-cyclopentyl)-hept-5-enoic acid methyl ester

(Z)-7-((1R,2S,3R,5R)-2-Butylcarbamoyloxymethyl-5-chloro-3-hydroxy-cyclopentyl)-hept-5-enoic acid

(Z)-7-((1R,2S,3R)-3-Hydroxy-5-oxo-2-phenethylcarbamoyloxymethyl-cyclopentyl)-hept-5-enoic acid methyl ester

(Z)-7-((1R,2S,3R)-3-Hydroxy-5-oxo-2-phenethylcarbamoyloxymethyl-cyclopentyl)-hept-5-enoic acid

(Z)-7-((1R,2S,3R)-2-Butylthiocarbamoyloxymethyl-3-hydroxy-5-oxo-cyclopentyl)-hept-5-enoic acid methyl ester

(Z)-7-((1R,2S,3R)-2-Butylthiocarbamoyloxymethyl-3-hydroxy-5-oxo-cyclopentyl)-hept-5-enoic acid.

22. The compound of claim 13 wherein said compound is selected from the group consisting of (Z)-7-((1R,2S,3R)-2-Butylcarbamoyloxymethyl-3-hydroxy-5-oxo-cyclopentyl)-hept-5-enoic acid methyl ester

(Z)-7-((1R,2S,3R)-2-Butylcarbamoyloxymethyl-3-hydroxy-5-oxo-cyclopentyl)-hept-5-enoic acid

(Z)-7-((1R,2S,3R,5R)-2-Butylcarbamoyloxymethyl-5-chloro-3-hydroxy-cyclopentyl)-hept-5-enoic acid methyl ester

(Z)-7-((1R,2S,3R,5R)-2-Butylcarbamoyloxymethyl-5-chloro-3-hydroxy-cyclopentyl)-hept-5-enoic acid

(Z)-7-((1R,2S,3R)-3-Hydroxy-5-oxo-2-phenethylcarbamoyloxymethyl-cyclopentyl)-hept-5-enoic acid methyl ester

(Z)-7-((1R,2S,3R)-3-Hydroxy-5-oxo-2-phenethylcarbamoyloxymethyl-cyclopentyl)-hept-5-enoic acid

(Z)-7-((1R,2S,3R)-2-Butylthiocarbamoyloxymethyl-3-hydroxy-5-oxo-cyclopentyl)-hept-5-enoic acid methyl ester

(Z)-7-((1R,2S,3R)-2-Butylthiocarbamoyloxymethyl-3-hydroxy-5-oxo-cyclopentyl)-hept-5-enoic acid.